Amendments to the Claims

- 1. (Currently amended) A pullulan-free pullulan-free edible film composition comprising:
 - (a) an effective amount of a film forming agent; and
- (b) an effective amount of an antimicrobial agent wherein the antimicrobial agent comprises Magnolia Bark extract.
- 2. (Currently amended) The composition of claim 1 wherein the film forming agent comprises a mixture of a maltodextrine matodextrin, a filler, and a hydrocolloid.
- 3. (Currently amended) The composition of claim 2 wherein the maltodextrine matodextrin comprises about 5 wt.% to about 60 wt.% of the edible film.
- 4. (Currently amended) The composition of claim 2 wherein the maltodextrine matodextrin comprises about 20 wt.% to about 40 wt.% of the edible film.
- 5. (Original) The composition of claim 2 wherein the hydrocolloid comprises about 10 wt.% to about 50 wt.% of the edible film.
- 6. (Original) The composition of claim 2 wherein the hydrocolloid comprises about 20 wt.% to about 30 wt.% of the edible film.
- 7. (Original) The composition of claim 2 wherein the filler comprises about 5 wt.% to about 30 wt.% of the edible film.
- 8. (Original) The composition of claim 2 wherein the filler comprises about 15 wt.% to about 25 wt.% of the edible film.
- 9. (Currently amended) The composition of claim 2 wherein the hydrocolloid comprises a material selected from the group consisting of a natural gums gum, a biosynthetic gums gum, a natural seaweeds seaweed, a natural plant extrudates extrudate, a natural fiber extracts extract, a gelatins gelatin, a biosynthetic process starchs starch, a cellulosic materials material, an alginates alginate, pectin, and combinations thereof.

- 10. (Currently amended) The composition of <u>claim 2</u> <u>claim 9</u> wherein the <u>natural</u> gum <u>hydrocolloid</u> comprises a gum selected from the group consisting of natural seed gum, guar gum, locust gum, tara gum, gum arabic, ghatti gum, agar gum, and xanthan gum.
- 11. (Currently amended) The composition of <u>claim 2</u> claim 9 wherein the <u>alginate</u> <u>hydrocolloid</u> comprises sodium alginate or calcium alginate.
- 12. (Currently amended) The composition of <u>claim 2</u> <u>claim 9</u> wherein the <u>natural</u> <u>seaweed hydrocolloid</u> comprises a carrageenan.
- 13. (Currently amended) The composition of claim 2 wherein the filler comprises a food-grade bulk filler selected from the group consisting of microcrystalline cellulose, a cellulose polymer polymer, magnesium carbonate, calcium carbonate, ground limestone, a silicates silicate, clay, talc, titanium dioxide, a calcium phosphates phosphate, and combinations thereof.
- 14. (Currently amended) The composition of <u>claim 2</u> claim 13 wherein the <u>cellulose</u> polymer <u>filler</u> comprises wood.
- 15. (Currently amended) The composition of <u>claim 2</u> <u>claim 13</u> wherein the <u>silicate</u> filler comprises magnesium or aluminum silicate, or <u>combinations thereof</u>.
- 16. (Currently amended) The composition of <u>claim 2</u> <u>claim 13</u> wherein the <u>calcium</u> <u>phosphate filler</u> comprises mono-calcium phosphate, di-calcium phosphate, or tri-calcium phosphate, or combinations thereof.
- 17. (Currently amended) The composition of claim 1 wherein the Magnolia Bark Extract extract comprises about 1 wt% to about 10 wt% of the edible film.
- 18. (Currently amended) The composition of claim 1 wherein the Magnolia Bark Extract extract comprises about 8 wt% of the edible film.
- 19. (Currently amended) The composition of claim 1 wherein the Magnolia Bark Extract comprises about 5 wt% of the edible film.

- 20. (Currently amended) The composition of claim 1 wherein the Magnolia Bark Extract extract comprises at least one of Magnolol and honokoil.
- 21. (Original) The composition of claim 1 further comprising an effective amount of a medicament.
- 22. (Currently amended) The composition of claim 21 wherein the medicament comprises an oral cleansing or breath freshening compound selected from the group consisting of a pH control agents agent, inorganic components for tartar or caries control, a breath freshening agents agent, an anti-plaque/anti-gingivitis agents agent, a saliva stimulating agents agent, a pharmaceutical agents agent, a nutraceutical agents agent, a vitamins vitamin, a mineral mineral, and combinations thereof.
- 23. (Currently amended) The composition of <u>claim 21</u> <u>claim 22</u> wherein the pH <u>control agent medicament</u> comprises urea.
- 24. (Currently amended) The composition of <u>claim 21</u> <u>claim 22</u> wherein the <u>inorganic components for tartar or caries control medicament comprises</u> <u>comprises</u> phosphates or fluorides.
- 25. (Currently amended) The composition of <u>claim 21</u> claim 22 wherein the breath freshening agent agent <u>medicament</u> comprises zinc gluconate.
- 26. (Currently amended) The composition of <u>claim 21 claim 22</u> wherein the <u>anti-plaque/anti-gingivitis agent medicament</u> comprises cholorhexidene, CPC, or triclosan, <u>or combinations thereof.</u>
- 27. (Currently amended) The composition of <u>claim 21</u> <u>claim 22</u> wherein the <u>saliva</u> <u>stimulating agent</u> medicament comprises a food acid.
- 28. (Currently amended) The composition of claim 27 wherein the food acid comprises an acid selected from the group consisting of citric, lactic, maleic, succinic, ascorbic, adipic, fumaric, and tartaric acids, and combinations thereof.

- 29. (Original) The composition of claim 1 further comprising an effective amount of a softening agent.
- 30. (Original) The composition of claim 29 wherein the softening agent comprises about 0 wt% to about 20 wt % of the edible film.
- 31. (Original) The composition of claim 29 wherein the softening agent comprises about 2 wt% to about 10 wt% of the edible film.
- 32. (Currently amended) The composition of claim 29 wherein the softening agent comprises a plasticizer including a compound selected from the group consisting of sorbitol, glycerin, polyethylene glycol, propylene glycol, hydrogenated starch hydrolysates, corn syrup, and combinations thereof.
- 33. (Original) The composition of claim 1 further comprising an effective amount of a coloring agent.
- 34. (Original) The composition of claim 1 further comprising an effective amount of a flavoring agent.
- 35. (Original) The composition of claim 34 wherein the flavoring agent comprises about 0.1 wt% to about 20 wt % of the edible film.
- 36. (Original) The composition of claim 34 wherein the flavoring agent comprises about 10 wt% to about 15 wt% of the edible film.
- 37. (Currently amended) The composition of claim 34 wherein the flavoring agent comprises a material selected from the group consisting of essential oils, synthetic flavors, fruit essences, anise, flavor oils with germ killing properties, and mixtures combinations thereof.
- 38. (Currently amended) The composition of <u>claim 34</u> claim 37 wherein the <u>essential</u> oils flavoring agent comprises an oil selected from the group consisting of citrus oil,

peppermint oil, spearmint oil, mint oil, clove oil, oil of wintergreen, and combinations thereof.

- 39. (Currently amended) The composition of <u>claim 34</u> claim 37 wherein the <u>flavor</u> oils with germ killing properties <u>flavoring agent</u> comprise <u>a material selected from the group consisting of menthol</u>, eucalyptol, thymol, and combinations thereof.
- 40. (Original) The composition of claim 1 further comprising an effective amount of an emulsifying agent.
- 41. (Currently amended) The composition of claim 40 wherein the emulsifying agent comprises a material selected from the group consisting of lecithin, (C_{10} - C_{18}) fatty acids, mono and diacyl glycerides, ox bile extract, polyglycerol esters, polyethylene sorbitan esters, propolyene glycol, sorbitan monopalmitate, sorbitan monosterate, sorbitan tristerate, enzyme modified lecithin, hyroxylated lecithins, and combinations thereof.
- 42. (Currently amended) A method of oral cleansing by applying a pullulan-free edible film to the oral cavity, wherein the edible film comprises:
 - (a) an effective amount of a film forming agent; and
- (b) an effective amount of an antimicrobial agent wherein the antimicrobial agent comprises Magnolia Bark Extract extract.
- 43. (Currently amended) The method of claim 42 wherein the Magnolia Bark Extract extract comprises at least about 1 wt% of the edible film.
- 44. (Currently amended) The method of claim 42 wherein the Magnolia Bark Extract extract comprises about 5 wt% of the edible film.
- 45. (Currently amended) The method of claim 42 wherein the Magnolia Bark Extract extract comprises at least one of Magnolol and honokoil.
- 46. (Currently amended) The method of claim 42 wherein the film forming agent comprises a mixture of a <u>maltodextrin</u> maltodextrine, a filler, and a hydrocolloid.

- 47. (Currently amended) The method of claim 46 wherein the <u>maltodextrin</u> maltodextrine comprises about 5 wt.% to about 60 wt.% of the edible film.
- 48. (Original) The method of claim 46 wherein the hydrocolloid comprises about 10 wt.% to about 50 wt.% of the edible film.
- 49. (Original) The method of claim 46 wherein the filler comprises about 5 wt.% to about 30 wt.% of the edible film.
- 50. (Currently amended) The method of claim 46 wherein the hydrocolloid comprises a material selected from the group consisting of a natural gums gum, a biosynthetic gums gum, a natural seaweeds seaweed, a natural plant extrudates extrudate, a natural fiber extracts extract, a gelatins gelatin, a biosynthetic process starchs starch, a cellulosic materials material, an alginates alginate, pectin, and combinations thereof.
- 51. (Currently amended) The method of <u>claim 46</u> claim 50 wherein the <u>natural gum</u> <u>hydrocolloid</u> comprises a gum selected from the group consisting of natural seed gum, guar gum, locust gum, tara gum, gum arabic, ghatti gum, agar gum, and xanthan gum.
- 52. (Currently amended) The method of <u>claim 46</u> claim 50 wherein the <u>alginate</u> <u>hydrocolloid</u> comprises sodium alginate or calcium alginate.
- 53. (Currently amended) The method of <u>claim 46</u> claim 50 wherein the natural seaweed hydrocolloid comprises a carrageenan.
- 54. (Currently amended) The method of claim 46 wherein the filler comprises a food-grade bulk filler selected from the group consisting of microcrystalline cellulose, a cellulose <u>polymers</u> polymer, magnesium carbonate, calcium carbonate, ground limestone, a <u>silicates</u> silicate, clay, talc, titanium dioxide, a calcium <u>phosphates</u>-phosphate, and combinations thereof.
- 55. (Currently amended) The composition of <u>claim 46</u> claim 54 wherein the <u>cellulose polymer</u> filler comprises wood.

- 56. (Currently amended) The method of <u>claim 46</u> <u>claim 54</u> wherein the <u>silicate filler</u> comprises magnesium or aluminum silicate, or combinations thereof.
- 57. (Currently amended) The method of <u>claim 6</u> <u>claim 54</u> wherein the <u>calcium</u> <u>phosphate filler</u> comprises mono-calcium phosphate, di-calcium phosphate, or tri-calcium phosphate, or combinations thereof.
- 58. (Original) The method of claim 42 wherein the edible film further comprises one or more of a medicament, a softening agent, a coloring agent, a flavoring agent, and an emulsifying agent.
- 59. (Currently amended) The method of claim 42 wherein the edible film delivers at least about 0.1wt% Magnolia Bark Extract extract to the oral cavity.
- 60. (Currently amended) The method of claim 42 wherein the edible film delivers at least about 0.01wt% Magnolia Bark Extract extract to the oral cavity.
- 61. (Currently amended) The method of claim 42 wherein the edible film delivers at least about 0.005wt% Magnolia Bark Extract extract to the oral cavity.
- 62. (Currently amended) A method of making a pullulan-free edible film comprising:
- (a) forming an aqueous solution that includes a maltodextrin, a hydrocolloid, and a filler;
- (b) adding an effective amount of an antimicrobial agent to the aqueous solution, wherein the antimicrobial agent comprises Magnolia Bark Extract extract; and
 - (c) drying the aqueous solution to form a dry edible film.
- 63. (Currently amended) The method of claim 62 wherein adding an effective amount of an antimicrobial agent comprises adding sufficient Magnolia Bark Extract extract such that the dry edible film comprises at least about 1 wt% Magnolia Bark Extract extract.

- 64. (Original) The method of claim 62 wherein adding an anti-microbial agent comprises adding at least one of Magnolol and honokoil.
- 65. (Currently amended) The method of claim 62 wherein forming an aqueous solution comprises adding sufficient <u>maltodextrin</u> maltodextrine such that the dry edible film comprises about 5 wt.% to about 60 wt.% <u>maltodextrin</u> maltodextrine.
- 66. (Original) The method of claim 62 wherein forming an aqueous solution comprises adding sufficient hydrocolloid such that the dry edible film comprises about 10 wt.% to about 50 wt.% hydrocolloid.
- 67. (Original) The method of claim 62 wherein forming an aqueous solution comprises adding sufficient filler such that the dry edible film comprises about 5 wt.% to about 30 wt.% filler.
- 68. (Original) The method of claim 62 wherein forming an aqueous solution further comprises adding one or more of a medicament, a softening agent, a coloring agent, a flavoring agent, and an emulsifying agent.
- 69. (Original) The method of claim 62 further comprising heating the aqueous solution to a temperature of about 40°C to about 60°C prior to drying the aqueous solution.
- 70. (Currently amended) A treatment method for reducing the number or activity of bacteria in the oral cavity comprising the steps of:
- (a) providing an edible film composition comprising Magnolia Bark Extract extract in an amount sufficient to kill or deactivate oral bacteria; and
- (b) causing a person in need of the treatment to consume the edible film composition whereby the bacteria in the oral cavity of the person is reduced or inactivated by the treatment.